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Application#		==		<del></del>	Inventor Name
09194321	6284798	150	10/18/1999	GUANIDINE DERIVATIVES, METHODS OF PREPARING THEM AND THEIR USE AS DRUGS	AMTMANN, EBERHARD
06892346	4851435	150	08/01/1986	ANTIVITAL AND ANTITUMOR XANTHATE PHARMACEUTICAL COMPOSITIONS	AMTMANN , EBERHARD
08337887	Not Issued	161	11/14/1994	ANTIVIRAL AND ANTITUMOR PHARMACEUTICAL COMPOSITIONS	AMTMANN , EBERHARD
07781125	Not Issued	161	03/13/1992	ANTI-TUMOR AGENTS HAVING REDUCED TOXICITY ON THE BASIS OF CYTOSTATIC AGENTS AND XANTHOGENATES	AMTMANN , EBERHARD
09784618	Not Issued	071	02/15/2001	MEDICAMENT CONTAINING PLATINUM COMPLEX COMPOUNDS AND THE USE THEREOF	AMTMANN, EBERHARD
09815992	Not Issued	164	03/23/2001	NEW GUANIDINE DERIVATIVES, PROCESSES FOR PREPARING THEM AND THEIR USE AS PHARMACEUTICAL COMPOSITIONS	AMTMANN, EBERHARD
10220905	Not Issued	019		SPHINGOMYELINASE ENZYMES AND USES RELATING THERETO	AMTMANN, EBERHARD
<u>09815995</u>	6444706	150	03/23/2001	GUANIDINE DERIVATIVES, PROCESSES FOR PREPARING THEM AND THEIR USE AS PHARMACEUTICAL COMPOSITIONS	AMTMANN, EBERHARD

Inventor Search Completed: No Records to Display.

Search Another: Inventor	Last Name	First Name	\$10000000 <b>1</b>
Scaren Amounci : inventor	Amtmann	Eberhard	Search

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=> s thioplatin

1 THIOPLATIN L2

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2001:422765 CAPLUS AN

- Antitumoral activity of a sulphur-containing platinum complex with an ΤT acidic pH optimum
- Amtmann, Eberhard; Zoller, Margot; Wesch, Horst; Schilling, Gerhard
- Department D0600, German Cancer Research Centre, Heidelberg, 69120, Germany
- Cancer Chemother. Pharmacol. (2001), 47(6), 461-466 SO

CODEN: CCPHDZ; ISSN: 0344-5704

- PB Springer-Verlag
- DTJournal
- LΑ English
- CC 1 (Pharmacology)
- Platinum complexes are essential tools for cancer treatment despite their AΒ toxic side effects. Here we describe a new platinum complex with sulphurs as complexing atoms (thioplatin). Purpose: To demonstrate that the antitumoral activity of a new sulfur-contg. platinum compd. ( thioplatin) depends on a slightly acidic pH.Methods: Platinum uptake by tumor cells and interaction with DNA was detd. at slightly acidic or alk. pH. To demonstrate low in vivo toxicity the effects of thioplatin on body wt., blood urea nitrogen, white blood cell count and the histopathol. appearance of small intestines and kidneys were evaluated at doses that displayed antitumoral effects against human small-cell lung cancer and human colorectal cancer xenotransplants in nude mice. Results: The slightly acidic pH optimum of thioplatin was proven by the altered electrophoretic mobility of plasmid DNA, quantitation of the platinum content in the DNA of tumor cells and cytotoxicity studies. Thioplatin displayed antitumoral activity without severe side effects such as wt. loss, renal ischemia, destruction of villi in the small intestine or leukopenia as obsd. at comparable doses of cisplatin. Furthermore, probably due to its lipophilic nature, thioplatin was taken up readily even by cisplatin-resistant cells. In vivo studies with human tumor xenografts in nude mice showed a therapeutic index of thioplatin five to ten times higher than that of cisplatin.

RE.CNT 13 THERE ARE 13 CITED REFERENCES AVAILABLE FOR THIS RECORD

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---Logging off of STN---

Executing the logoff script...

=> LOG Y

SINCE FILE	TOTAL
ENTRY	SESSION
5.21	9.68
SINCE FILE	TOTAL
ENTRY	SESSION
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